This meeting is for the purpose of review, discussion, evaluation, and recommendations on financial assistance under the National Foundation on the Arts and the Humanities Act of 1965, as amended, including information given in confidence to the agency. In accordance with the determination of the Chairman of May 22, 2001, these sessions will be closed to the public pursuant to subsection (c)(4), (6) and (9)(B) of section 552b of Title 5, United States Code.

Further information with reference to this meeting can be obtained from Ms. Kathy Plowitz-Worden, Panel Coordinator, National Endowment for the Arts, Washington, DC, 20506, or call 202/682–5691.

Dated: June 21, 2001.

Kathy Plowitz-Worden,

Panel Coordinator, Panel Operations, National Endowment for the Arts.

[FR Doc. 01–16087 Filed 6–26–01; 8:45 am]

BILLING CODE 7537–01–P

NORTHEAST DAIRY COMPACT COMMISSION

Notice of Meeting

AGENCY: Northeast Dairy Compact Commission.

ACTION: Notice of meeting.

SUMMARY: The Compact Commission will hold its regular monthly meeting to consider matters relating to administration and enforcement of the price regulation. This meeting will be held in So. Portland, Maine, continuing the Commission's program of holding a meeting in each of the Compact states. In addition to receiving reports and recommendations of its standing Committees, the Commission will receive a number of informational reports about the impact of the over-order price regulation in Maine.

DATES: The meeting will begin at 10 a.m. on Monday, July 9, 2001.

ADDRESSES: The meeting will be held at the Best Western Merry Manor, 700 Main Street, So. Portland, Maine 04106.

FOR FURTHER INFORMATION CONTACT: Daniel Smith, Executive Director, Northeast Dairy Compact Commission, 64 Main Street, Room 21, Montpelier, VT 05602. Telephone (802) 229–1941.

Authority: 7 U.S.C. 7256.

Dated: June 20, 2001.

Daniel Smith,

Executive Director.

[FR Doc. 01–16044 Filed 6–26–01; 8:45 am]

BILLING CODE 1650–01–P

NUCLEAR REGULATORY COMMISSION

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: U. S. Nuclear Regulatory Commission (NRC).

ACTION: Notice of pending NRC action to submit an information collection request to OMB and solicitation of public comment.

SUMMARY: The NRC is preparing a submittal to OMB for review of continued approval of information collections under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35).

Information pertaining to the requirement to be submitted:

1. *The title of the information collection:* 10 CFR 35.32 and 35.33 ''Quality Management Program and Misadministrations''.

2. *Current OMB approval number:* 3150–0171.

3. *How often the collection is required:* For quality management program (QMP): *Reporting:* New applicants for medical use licenses, who plan to use byproduct material in limited diagnostic and therapy quantities under Part 35, must develop a written QMP and submit a copy of it to NRC. When a new modality involving therapeutic quantities of byproduct material is added to an existing license, current licensees must submit QMP modifications. This ICR burden estimate is inflated by the one-time cost for the development and submission of QMPs for approximately 2000 Agreement States licensees in the ten Agreement States who have not adopted the rule and are not required to. *Recordkeeping:* Records of written directives, administered dose or dosage, annual review, and recordable events, for 3 years.

For Misadministrations: Reporting: Whenever a misadministration occurs. *Recordkeeping:* Records of misadministrations for 5 years.

4. *Who is required or asked to report:* NRC Part 35 licensees who use byproduct material in limited diagnostic and therapeutic ranges and similar type of licensees regulated by Agreement States.

5. *The estimated number of annual respondents:* 6300 (for both reporting and recordkeeping).

6. *The number of hours needed annually to complete the requirement or request:* 34,743 hours for applicable licensees (Reporting: 24,400 Hrs/yr, and Recordkeeping: 10,343 Hrs/yr, or an average of 5.5 hrs per licensee).

7. *Abstract:* In the medical use of byproduct material, there have been instances where byproduct material was not administered as intended or was administered to a wrong individual, which resulted in unnecessary exposures or inadequate diagnostic or therapeutic procedures. The most frequent causes of these incidents were: insufficient supervision, deficient procedures, failure to follow procedures, and inattention to detail. In an effort to reduce the frequency of such events, the NRC requires licensees to implement a quality management program (§ 35.32) to provide high confidence that byproduct material or radiation from byproduct material will be administered as directed by an authorized user physician. Collection of this information enables the NRC to ascertain whether misadministrations (§ 35.33) are investigated by the licensee and that corrective action is taken. Additionally, NRC has a responsibility to inform the medical community of generic issues identified in the NRC review of misadministrations.

Revisions to 10 CFR 35.32 and 35.33 are being made as part of a complete revision of 10 CFR Part 35 to incorporate specific improvements in NRC's regulations governing the medical use of byproduct material. A final rule revising Part 35 was affirmed by the Commission on October 23, 2000 and was submitted, along with its associated clearance package, to the Office of Management and Budget (OMB). A notice was published in the **Federal Register** on March 16, 2001, announcing a 30-day public comment period on the submittal. It is anticipated that the effective date of the final rule revising Part 35, including the revisions to Sections 35.32 and 35.33, will be March 2002, and the OMB clearance for Sections 35.32 and 35.33 will be then be included under the OMB clearance for Part 35 (3150–0010).

Currently, the OMB clearances for Sections 35.32 and 35.33 are due to expire October 31, 2001. In view of the fact that these parts will shortly thereafter be covered under OMB clearance 3150–0010, the Commission is seeking a 1-year clearance extension for the information collection requirments in these sections to allow sufficient time for OMB to complete its review of the NRC clearance package for the revision to Part 35, for NRC to publish the final rule, and for the rule to become effective. Because the final Part 35 and its OMB clearance will be in place in a short time period, the burden hour estimates in this extension package are not being revised from those contained in the previous OMB approval for